

**PCT**WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>5</sup> :</b> <b>C09J 7/02, 133/08</b>	<b>AI</b>	<b>(11) International Publication Number:</b> <b>WO 92/13924</b> <b>(43) International Publication Date:</b> 20 August 1992 (20.08.92)
<b>(21) International Application Number:</b> PCT/US92/00613 <b>(22) International Filing Date:</b> 24 January 1992 (24.01.92) <b>(30) Priority data:</b> 651,468 6 February 1991 (06.02.91) US <b>(71) Applicant:</b> MINNESOTA MINING AND MANUFACTURING COMPANY [US/US]; 3M Center, Post Office Box 33427, Saint Paul, MN 55133-3427 (US). <b>(72) Inventors:</b> STEELMAN, Ronald, S. ; Post Office Box 33427, Saint Paul, MN 55133-3427 (US). CRANDALL, Michael, D. ; Post Office Box 33427, Saint Paul, MN 55133-34327 (US). DELGADO, Joaquin ; Post Office Box 33427, Saint Paul, MN 55133-3427 (US).		<b>(74) Agents:</b> ODAR, Thomas, J. et al.; Intellectual Property Counsel, Minnesota Mining and Manufacturing Company, Post Office Box 33427, Saint Paul, MN 55133-3427 (US). <b>(81) Designated States:</b> AT (European patent), AU, BE (European patent), BR, CA, CH (European patent), CS, DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), HU, IT (European patent), JP, KR, LU (European patent), MC (European patent), NL (European patent), SE (European patent). <b>Published</b> <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
<b>(54) Title:</b> POSITIONABLE ADHESIVE SYSTEM WITH HIGH SHEAR STRENGTH  <b>(57) Abstract</b>  A positionable aqueous adhesive composition is provided, comprising a blend or mixture of (a) an aqueous suspension of hollow, polymeric, inherently tacky elastomeric microspheres, and (b) an aqueous film-forming pressure sensitive adhesive latex. The adhesive is positionable, in that it allows for positional adjustment after application to a receptor, yet displays high peel adhesion and shear strength.		

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FI	Finland	ML	Mali
AU	Australia	FR	France	MN	Mongolia
BB	Barbados	GA	Gabon	MR	Mauritania
BE	Belgium	GB	United Kingdom	MW	Malawi
BF	Burkina Faso	GN	Guinea	NL	Netherlands
BG	Bulgaria	GR	Greece	NO	Norway
BJ	Benin	HU	Hungary	PL	Poland
BR	Brazil	IE	Ireland	RO	Romania
CA	Canada	IT	Italy	RU	Russian Federation
CF	Central African Republic	JP	Japan	SD	Sudan
CG	Congo	KP	Democratic People's Republic of Korea	SE	Sweden
CH	Switzerland	KR	Republic of Korea	SN	Senegal
CI	Côte d'Ivoire	LJ	Liechtenstein	SU	Soviet Union
CM	Cameroon	LK	Sri Lanka	TD	Chad
CS	Czechoslovakia	LU	Luxembourg	TG	Togo
DE	Germany	MC	Monaco	US	United States of America
DK	Denmark	MG	Madagascar		
ES	Spain				

- 1 -

**POSITIONABLE ADHESIVE SYSTEM WITH HIGH SHEAR STRENGTH**Field of the Invention

This invention relates to a positionable  
5 adhesive system displaying final elevated shear  
strength and peel adhesion, comprising a mixture of  
hollow, polymeric, acrylate, infusible, inherently  
tacky elastomeric microspheres together with a  
polymeric film-forming pressure sensitive latex  
10 adhesive.

Background of the InventionDescription of the Related Art

15 Positionable adhesives are those which allow  
for placement of an article containing such an adhesive  
onto a receptor in an exact position, because the  
article can be adjusted relative to the receptor after  
initial placement.

20 In some instances, the adhesive can be  
designated repositionable or repeatedly reuseable.  
Such adhesives exhibit aggressive tack but low peel  
adhesion properties, thus allowing repeated  
reuseability. Commercial products such as the 3M Brand  
25 Post-it<sup>TM</sup> Notes display such adhesive characteristics.

Positionable adhesive systems, however, as used  
herein, generally display high peel adhesion and  
elevated shear strength after positioned placement.

Such systems have been disclosed for utility  
30 such as in the creation of sign faces or other graphic  
systems utilizing a vinyl film. For example, U.S.  
Patent Nos. 3,314,838; 3,331,729 and 3,413,168 disclose  
the concept of positionability based on a conventional  
film-forming adhesive having contained therein  
35 frangible glass bubbles. Positionability is obtained  
by the fact that the glass bubbles, during initial  
application of the coated article,

- 2 -

allow for incomplete surface contact of the adhesive to the ultimate receptor. Once the article is properly positioned, pressure on the face of the article causes a rupture of the glass bubbles, thus allowing the full surface area to be contacted with the receptor, forming a strong bond. Obviously, once the frangible bubbles are ruptured, any further positionability characteristics are lost. In addition, articles coated with the adhesive containing the frangible bubbles inherently have added manufacturing steps and furthermore require special liners for application.

Reported adhesive systems for repositionable products range from the microsphere adhesives disclosed in U.S. Patent Nos. 3,691,140 and 4,166,152 to pattern-coated conventional adhesive systems. A further definition of an adhesive system containing such microspheres is disclosed in U.S. Patent No. 3,857,731, wherein a binder material is mixed with the microspheres. In some examples in this latter patent, the binder is an acrylate pressure sensitive adhesive. However, in all instances, the adhesion characteristics of the microspheres themselves are not taught to be overshadowed by the pressure sensitive adhesive. In this instance, the binder creates a physical "socket" to mechanically hold the individual microspheres onto a substrate, thus asserting to prevent undesired transfer of the microspheres to a receptor, increasing the retention of desired adhesive features.

U.S. Patent No. 4,735,837 discloses a similar system, wherein solid microspheres are mixed with a binder resin, which is taught to be a pressure sensitive adhesive. In this disclosure, the characteristics of adhesion and repositionability come from the pressure sensitive adhesive rather than the microspheres themselves. In fact, the patent discloses that the microspheres need not have adhesion capabilities at all.

- 3 -

In EPO 209,337, it is disclosed that the basic deficiency of solid microspher adhesives is that they exhibit excessive transfer. In this instance, a reactive monomer is included with the adhesive  
5 microspheres during their preparation. The monomer remains unreacted during the adhesive polymerization, but subsequently undergoes a reaction with the binder and/or the backing to provide shear values of up to 50 Newtons, claimed to be higher than other  
10 microsphere-based systems.

In contrast with the foregoing, we have now discovered that an aqueous pressure sensitive adhesive system can be made utilizing a combination of hollow microspheres which are polymeric, acrylate, infusible  
15 and inherently tacky, together with a polymeric film-forming pressure sensitive adhesive. This aqueous blend, when applied to a backing, displays the desired positionability, ease of application, and typically associated with the positionable adhesives discussed  
20 above, and in addition displays the high shear strength, high peel adhesion and other performance characteristics normally associated with conventional film-forming pressure sensitive adhesives. In addition, we have found that this system provides for  
25 clean removal from most receptors.

#### Summary of the Invention

In accordance with the invention there is  
30 provided a positionable aqueous adhesive composition and article containing same thereon, the composition comprising a blend or mixture of (a) an aqueous suspension of hollow, polymeric, infusible, inherently tacky elastomeric microspheres, and (b) an aqueous  
35 film-forming pressure sensitive adhesive latex comprising at least one long chain alkyl acrylate. Preferably, the weight ratio of microspheres to latex,

- 4 -

based on solids, is from about 12 to 1 to about 39 to 1.

The adhesive is positionable, i.e., can be adjusted positionally after application to a receptor, yet displays high peel adhesion and shear strength, and displays clean removal from most receptors.

Graphic or decorative films, e.g., polyvinyl chloride can be applied to a receptor and positioned accurately, yet final peel adhesion and shear forces will be sufficiently elevated to allow the film to remain in place without movement or deformation.

#### Detailed Description of the Invention

The hollow polymeric microspheres having utility herein are disclosed in detail in commonly assigned copending application Serial No. 276,767, incorporated herein by reference. Such microspheres comprise at least about 85 parts by weight of at least one alkyl acrylate or alkyl methacrylate ester together with up to about 15 parts by weight of at least one polar monomer, a majority of such microspheres having one or more interior voids. Preferably, the microspheres contain a central cavity at least 10%, and most preferably, 30% of the diameter of the microsphere itself. An aqueous suspension of such hollow microspheres can be prepared by emulsification processes as disclosed in the subject application. Basically, they can be prepared by a two-step process comprised of forming a water-in-oil emulsion of an aqueous solution of one or more polar monomers in oil phase monomers; forming a water-in-oil-in-water emulsion by dispersing the first emulsion into an aqueous phase; followed by initiating polymerization, preferably by the application of heat or radiation. In addition, aqueous suspensions of such microspheres which contain moderately ionized polar monomers may be prepared by a simpler one-step process comprising the aqueous suspension polymerization of at least one alkyl

- 5 -

acrylate or methacrylate ester monomer and at least one n-ionic polar monomer in the presence of at least one emulsifier capable of producing a water-in-oil emulsion inside the droplets, which is substantially stable during polymerization. Both methods produce an aqueous suspension of monomer droplets which upon polymerization become microspheres, the majority of which contain at least one interior cavity as discussed above.

10           The aqueous pressure sensitive adhesive latex used to blend with the aqueous suspension of hollow microspheres is composed of at least one long chain alkyl acrylate, i.e., containing from 4 to 12 carbon atoms, and preferably a polar monomer.

15           The film-forming pressure sensitive adhesive should be used at a level lower than the interspatial volume of the hollow microspheres. Preferably, the adhesive mixture contains a microsphere to film-forming adhesive ratio of from 39:1 to 12:1. Levels of  
20 film-forming adhesive significantly higher than this may produce adhesive coated articles that retain the characteristics of the film-forming pressure sensitive adhesive, detrimental to positionability.

          The aqueous composition can be conventionally  
25 applied onto materials such as paper, and vinyl and polyester films, as by knife or notched-bar coating techniques, for example.

          Additives can be included in the mixture to enhance specific objectives, such as the coating of the  
30 aqueous system or modifying performance characteristics thereof.

          Anti-foams can be added, especially when the aqueous system of the invention is to be applied as a coating at high rates of speed. One example of such a  
35 material is "Foamaster" JMY, a conventional anti-foam agent from Henkel Process Chemicals, Inc.

- 6 -

Colorants can be added to enhance visual appearance, quality control, etc. Several vendors provide concentrates which add such desirable features.

Chemical crosslinkers, such as polyaziridines, can increase ply adhesion, high temperature shear and cohesive strengths. Such improvements, however, typically can provide a loss in tack and ultimate adhesion.

Viscosity control materials can be added to assist in coating quality and control. Examples include water soluble materials such as various cellulose products, polyacrylic acids, and vinyl alcohols.

The invention will now be described by the use of the following non-limiting examples, wherein all parts are by weight unless otherwise specified.

#### Example 1

Hollow microspheres are manufactured in accordance with the following procedure:

In a one-liter resin reactor equipped with mechanical stirrer, thermometer and inlet-outlet lines for vacuum and nitrogen, 450 grams of deionized water, 141 grams of iso-octyl acrylate, 9.0 grams of acrylic acid and 0.5 gram of benzoyl peroxide were charged. Vacuum was applied to evacuate the reactor atmosphere and the reactor was purged with argon. The agitation was set to 400 RPM and when the initiator had dissolved, 1.5 grams of ammonium lauryl sulfate (Standapol A, from Henkel AG) were added. The temperature of the reactor was raised to 60°C and maintained for 22 hours, an argon purge being maintained during the polymerization. The suspension was then allowed to cool to room temperature. The reactor was then emptied and the suspension filtered. Optical microscopy showed hollow microspheres from about 4 to about 90 microns in diameter, the majority



- 7 -

of the microspheres containing a central cavity of at least 30% of the diameter of the microspheres.

This microsphere suspension exhibits a pH typically between 2.0 and 4.0 and has a solids content of approximately 25%. With this procedure, the mean sphere diameter is typically 40-60 microns. (Different vessel proportions or mixing rates will alter the particle size, void diameter, etc.)

An aqueous pressure sensitive adhesive was prepared as follows:

A split-resin flask of 2000 ml. capacity was fitted with a variable speed agitator, condenser, purging tube for introducing nitrogen, and a recording controller. The following materials were added to the flask while purging same with nitrogen:

404 grams	Distilled water
1.50 grams	Sodium dodecylbenzene sulfonate
435 grams	Isooctyl acrylate
60 grams	N-tert-octylacrylamide
0.60 gram	Sodium bicarbonate
5.0 grams	Sodium styrene sulfonate

The solid N-tert-octylacrylamide was dissolved in the isooctyl acrylate before adding to the flask. The nitrogen purge was continued until the end of the run. The flask and its contents were heated to 50°C, at which temperature an initiator charge of 0.05 gram potassium persulfate and 0.0125 gram sodium metabisulfite was added. The reaction temperature was held at 50°C for approximately 24 hours to complete the polymerization. The resulting latex had no coagulum and the solids content was 54%.

The adhesives as prepared above were combined by adding, to a suitable container, 95.88 parts of the aqueous microsphere suspension at 25% solids; 3.6 parts of the aqueous pressure sensitive adhesive containing 54% solids; 0.52 part of ASE-60, commercially available

- 8 -

from Rohm & Haas, as a rheological control agent. The pH was then raised with ammonium hydroxide to between 7.5 and 8.5.

The composition was then knife coated onto a polyvinyl chloride film and dried at 200°F for 1 minute to provide a dry coating weight of 19 grams per square meter.

#### Example 2

The adhesive of Example 1 was duplicated with the exception of utilizing 98.77 parts of the microsphere suspension and 1.23 parts of the pressure sensitive adhesive latex. This was applied to a polyvinyl chloride film as per Example 1.

#### Example 3

An adhesive blend was prepared by mixing, in a suitable container, 96.23 parts of the aqueous microsphere suspension of Example 1; 2.99 parts of Gelva 2397, an aqueous pressure sensitive adhesive available from Monsanto, containing 65% solids; 0.52 part of ASE-60; followed by raising the pH with ammonium hydroxide to between 7.5 and 8.5; and the addition of 0.26 part of C-E2 color concentrate from Ciba-Geigy to provide opacity.

The composition was coated onto a polyvinyl chloride film at a dry coating weight of 19 grams per square meter, as per Example 1.

#### Example 4

A film-forming pressure sensitive adhesive was manufactured as follows:

A split-resin flask of 2000 ml. capacity was fitted with a variable speed agitator, condenser, purging tube for introducing nitrogen, and a recording controller. The following materials were added to the flask while purging the flask with nitrogen:

600 grams	Distilled water
-----------	-----------------

- 9 -

	4.80 grams	Sodium dodecylbenzene sulfonate
	4.80 grams	Nonyl phenol - 10.5 mole ethylene oxide adduct
	16.0 grams	Acrylic acid
5	160.0 grams	Ethyl acrylate
	160.0 grams	Isooctyl acrylate
	64.0 grams	Butyl acrylate

The nitrogen purge was continued until the end of the  
10 run. The flask and its contents were agitated at 300  
rpm, and was heated to 32°C. At this temperature an  
initiator charge of 0.30 gram potassium persulfate and  
0.08 gram sodium metabisulfite was added. An  
exothermic reaction occurred increasing the temperature  
15 to approximately 75°C after which the reaction mixture  
was allowed to cool. The resulting latex had no  
coagulum and the solids content was 40%.

An adhesive blend was then made by mixing 97.68  
parts of the aqueous microsphere suspension of Example  
20 1 with 1.57 parts of the film-forming pressure  
sensitive adhesive, together with 0.5 part of ASE-60,  
adjustment of pH to 7.5 to 8.5 with ammonium hydroxide,  
and the addition of 0.25 part of C-E2 color  
concentrate.

25 The blend was coated as per Example 1 onto a  
polyvinyl chloride film at a coating weight of 19 grams  
per square meter.

#### Example 5

30 A film-forming pressure sensitive adhesive was  
made in accordance with Example 4 with the exception  
that the monomer mixture was comprised of 320 grams of  
ethyl acrylate, 64 grams of butyl acrylate, and 16  
grams of acrylic acid. A blend was then prepared by  
35 adding 94.46 parts of the aqueous microsphere  
suspension of Example 1 to 4.77 parts of the  
film-forming pressure sensitive adhesive, again  
containing 40% solids, 0.52 part of ASE-60, pH

- 10 -

adjustment to 7.5 to 8.5 with ammonium hydroxide, and the addition of 0.26 part of C-E2 color concentrate.

Again, the blend was applied onto polyvinyl chloride film at a coating weight of 19 grams per 5 square centimeter as per Example 1.

The foregoing samples were tested for adhesion, both initially and after aging, and dynamic shear was also determined. These results are reported in TABLE 1.

10 For the adhesion tests, a 25.4 mm strip of sample was applied to a test panel using three passes with a 2.27 kg roller. The adhesion was tested by peeling the adhered sample at 180 degrees at a rate of 2.57 mm per minute. The value reported is an average 15 reading over a five second period.

Samples were also aged at room temperature (27°C) for the designated time.

For the dynamic shear test, a .076 mm polyester tape was laminated to a 26.5 mm wide sample to be 20 tested. The samples were adhered to a test panel, typically a painted metal panel, and trimmed such that a 25.7 mm by 25.7 mm contact area was retained. The sample was pulled with an Instron testing machine at a rate of 5 mm/minute. The maximum force required to 25 remove the sample from the test panel is recorded.

TABLE 1

30	Initial	Adhesion	
	Adhesion,	After 24 Hours,	Dynamic shear,
	Newtons Per	Newtons per	Newtons Per
	Decimeter	Decimeter	Decimeter
	<u>Width</u>	<u>Width</u>	<u>Square</u>
35 Example 1	20.84	27.50	1176.92
Example 2	22.07	20.97	1117.99

- 11 -

Example 3	23.67	26.45	1228.06
Example 4	20.67	26.09	993.18
5 Example 5	27.34	27.32	1037.52

Visual observation of the samples showed no  
sign of adhesive transfer to the receptor test  
10 specimens.

For best performance, the blend should be  
applied at the minimum coating thickness required to  
obtain a monolayer of microspheres. High coating  
thicknesses are basically wasteful, while reduced  
15 coating thicknesses can result in incomplete surface  
coverage, and thus reduced peel adhesion and shear  
values. The preferred coating thickness is represented  
by the corresponding diameter of the average  
microsphere. For example, microspheres made by the  
20 aforementioned procedure typically average 40 to 60  
microns in diameter, and thus the preferred coating  
thickness would be in that same range.

- 12 -

In the Claims

1. An aqueous adhesive composition capable of displaying positionablity, when applied as a layer to a  
5 substrate, while maintaining elevated shear strength and peel adhesion, comprising an aqueous blend of
- (a) an aqueous suspension of hollow, polymeric, infusible, inherently tacky elastomeric microspheres; and
  - 10 (b) an aqueous film-forming pressure sensitive adhesive latex comprising at least one long chain alkyl acrylate.

2. The composition of claim 1 wherein said  
15 microspheres are comprised of at least about 85 parts by weight of at least one alkyl acrylate or methacrylate ester and up to about 15 parts by weight of at least one polar monomer.

20 3. The composition of claim 1 wherein the weight ratio of said microspheres to said latex, on a solids basis, is from about 12 to 1 to about 39 to 1.

4. The composition of claim 1 wherein said  
25 latex further comprises at least one polar monomer.

5. An article comprising a substrate, having applied to at least a portion of at least one surface thereof, a coating of a positionable adhesive  
30 composition comprising

- (a) an aqueous suspension of hollow, polymeric, infusible, inherently tacky elastomeric microspheres; and
- (b) an aqueous film-forming pressure sensitive  
35 adhesive latex comprising at least one long chain alkyl acrylate.

6. The article of claim 5 wherein said substrate is a polyvinyl chloride film.

- 13 -

7. The article of claim 5 wherein said microspheres are comprised of at least about 85 parts by weight of at least one alkyl acrylate or methacrylate ester and up to about 15 parts by weight of at least one polar monomer.

8. The article of claim 5 wherein the weight ratio of said microspheres to said latex, on a solids basis, is from about 12 to 1 to about 39 to 1.

10

9. The article of claim 5 wherein said latex further comprises at least one polar monomer.

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 92/00613

<b>I. CLASSIFICATION OF SUBJECT MATTER</b> (If several classification symbols apply, indicate all) <sup>6</sup>		
According to International Patent Classification (IPC) or to both National Classification and IPC Int.Cl. 5 C09J7/02; C09J133/08		
<b>II. FIELDS SEARCHED</b>		
Minimum Documentation Searched <sup>7</sup>		
Classification System	Classification Symbols	
Int.Cl. 5	C09J	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched <sup>8</sup>		
<b>III. DOCUMENTS CONSIDERED TO BE RELEVANT<sup>9</sup></b>		
Category *	Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup>	Relevant to Claim No. <sup>13</sup>
Y	US,A,4 968 562 (J.DELGADO) 6 November 1990 see column 5, line 44 - line 49 see column 8, line 1 - line 7; claims 1-11 ---	1-5
Y,P	EP,A,0 419 020 (3M) see claims 1-4,6,9,10 ---	1-5
<p>* Special categories of cited documents: <sup>10</sup></p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"A" document member of the same patent family</p>		
<b>IV. CERTIFICATION</b>		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
15 JUNE 1992	10. 07 92	
International Searching Authority	Signature of Authorized Officer	
EUR PEAN PATENT OFFICE	BUSCAGLIONE Y. <i>[Signature]</i>	



**ANNEX TO THE INTERNATIONAL SEARCH REPORT  
ON INTERNATIONAL PATENT APPLICATION NO. US 9200613  
SA 57768**

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on  
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information. 15/06/92

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US-A-4968562	06-11-90	AU-A- 6928391	29-08-91
		EP-A- 0444354	04-09-91
		US-A- 4988567	29-01-91
EP-A-0419020	27-03-91	US-A- 4994322	19-02-91
		AU-A- 5989690	21-03-91
		CA-A- 2021958	19-03-91
		JP-A- 3111473	13-05-91

EPO FORM P0079

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

